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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.		
10/813,177	03/29/2004	Wei Gu	5199-178	5834		
Leslie Gladstor	7590 10/17/2007 ne Restaino	EXAM	EXAMINER			
Brown Raysman Millstein Felder & Steiner LLP 163 Madison Avenue P.O. Box 1989 Morristown, NJ 07962-1989			FETTEROLF,	FETTEROLF, BRANDON J		
			ART UNIT	PAPER NUMBER		
			1642			
			MAIL DATE	DELIVERY MODE		
			10/17/2007	PAPER		

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

# Advisory Action Before the Filing of an Appeal Brief

Application No.	Applicant(s)		
10/813,177	GU ET AL.		
Examiner	Art Unit		
Brandon J. Fetterolf, PhD	1642		

	Brandon J. Fetterolf, PhD	1642				
The MAILING DATE of this communication appear	ars on the cover sheet with the c	orrespondence add	ress			
THE REPLY FILED 31 August 2007 FAILS TO PLACE THIS APPLICATION IN CONDITION FOR ALLOWANCE.						
<ul> <li>1.  The reply was filed after a final rejection, but prior to or on this application, applicant must timely file one of the follow places the application in condition for allowance; (2) a No a Request for Continued Examination (RCE) in compliance time periods:</li> <li>a) The period for reply expires 3 months from the mailing date</li> </ul>	the same day as filing a Notice of ving replies: (1) an amendment, aff tice of Appeal (with appeal fee) in ce with 37 CFR 1.114. The reply must	Appeal. To avoid aba idavit, or other evider compliance with 37 C	nce, which FR 41.31; or (3)			
b) The period for reply expires on: (1) the mailing date of this A no event, however, will the statutory period for reply expire la Examiner Note: If box 1 is checked, check either box (a) or (TWO MONTHS OF THE FINAL REJECTION. See MPEP 76	dvisory Action, or (2) the date set forth ater than SIX MONTHS from the mailing b). ONLY CHECK BOX (b) WHEN THE 06.07(f).	g date of the final rejecti E FIRST REPLY WAS F	on. ILED WITHIN			
Extensions of time may be obtained under 37 CFR 1.136(a). The date have been filed is the date for purposes of determining the period of extunder 37 CFR 1.17(a) is calculated from: (1) the expiration date of the set forth in (b) above, if checked. Any reply received by the Office later may reduce any earned patent term adjustment. See 37 CFR 1.704(b) NOTICE OF APPEAL	ension and the corresponding amount shortened statutory period for reply origing than three months after the mailing da	of the fee. The appropr inally set in the final Offi te of the final rejection,	iate extension fee ce action; or (2) as even if timely filed,			
2. The Notice of Appeal was filed on A brief in comp filing the Notice of Appeal (37 CFR 41.37(a)), or any extension a Notice of Appeal has been filed, any reply must be filed AMENDMENTS	nsion thereof (37 CFR 41.37(e)), to	avoid dismissal of th	ns of the date of ne appeal. Since			
3. The proposed amendment(s) filed after a final rejection,	but prior to the date of filing a brief	. will not be entered b	ecause			
<ul> <li>(a)    ☐ They raise new issues that would require further co</li> <li>(b) ☐ They raise the issue of new matter (see NOTE below)</li> </ul>	nsideration and/or search (see NO w);	TE below);				
(c) They are not deemed to place the application in befappeal; and/or	ter torm for appeal by materially re	aucing or simplifying	the issues for			
(d) They present additional claims without canceling a	corresponding number of finally rej	ected claims.				
NOTE: (See 37 CFR 1.116 and 41.33(a)).						
4. The amendments are not in compliance with 37 CFR 1.1		ompliant Amendment	(PTOL-324).			
5. Applicant's reply has overcome the following rejection(s)		time also filed amounders	ant concelling the			
6. Newly proposed or amended claim(s) would be all non-allowable claim(s).						
7. For purposes of appeal, the proposed amendment(s): a) how the new or amended claims would be rejected is pro The status of the claim(s) is (or will be) as follows:	☑ will not be entered, or b) ☐ wi vided below or appended.	II be entered and an	explanation of			
Claim(s) allowed: Claim(s) objected to:						
Claim(s) rejected: <u>47,48,54,57 and 61-63</u> . Claim(s) withdrawn from consideration:						
AFFIDAVIT OR OTHER EVIDENCE	it hafara ar an tha data of filing a N	otice of Anneal will n	nt he entered			
<ol> <li>The affidavit or other evidence filed after a final action, but because applicant failed to provide a showing of good an was not earlier presented. See 37 CFR 1.116(e).</li> </ol>	d sufficient reasons why the affida	vit or other evidence i	s necessary and			
9. The affidavit or other evidence filed after the date of filing entered because the affidavit or other evidence failed to o showing a good and sufficient reasons why it is necessar	overcome <u>all</u> rejections under appe y and was not earlier presented. S	al and/or appellant fa See 37 CFR 41.33(d)(	ils to provide a 1).			
10.  The affidavit or other evidence is entered. An explanation REQUEST FOR RECONSIDERATION/OTHER	n of the status of the claims after e	entry is below or attac	hed.			
11. The request for reconsideration has been considered by	it does NOT place the application i	n condition for allowa	nce because:			
<ul> <li>12. ☑ Note the attached Information Disclosure Statement(s).</li> <li>13. ☐ Other:</li> </ul>	(PTO/SB/08) Paper No(s). <u>1/7/200</u>	<u>5</u>				
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#### **DETAILED ACTION**

### Response to the Amendment

The Amendment filed on 8/31/2007 in response to the previous Final Office Action (5/31/2007) is acknowledged, but has not been entered. The amendment has not been entered because it contains claim limitations, which were not previously searched and/or considered. For example, Claim 54 is drawn to a method for determining whether an agent is reactive with Mdm2 comprising the steps of contacting a candidate agent with Mdm2, in the presence of herpesvirus-associated ubiquitin-specific protease (HASUP); and (b) determining whether the candidate agent inhibits Mdm2-HAUSP protein complex formation, wherein inhibition of Mdm2-HAUSP protein complex formation indicates that the agent is reactive with Mdm2. Similarly, Claim 57 recites a method for determining whether an agent is reactive with HAUSP comprising the steps of contacting a candidate agent with HAUSP, in the presence of Mdm2; and (b) determining whether the candidate agent inhibits Mdm2-HAUSP protein complex formation, wherein inhibition of Mdm2-HAUSP protein complex formation indicates that the agent is reactive with HAUSP. Thus, the claims appear to encompass determining the same Mdm2-HAUSP protein complex for the determination of an active agent against two distinct proteins; and therefore, would require further consideration with respect to 112, 2<sup>nd</sup> paragraph, as well as 112, 1<sup>st</sup> paragraph enablement.

Claims 47-48, 54, 57, 61-63 are currently pending and under consideration.

#### Information Disclosure Statement

The Information Disclosure Statement filed 1/7/2005 is acknowledged. The submission is in compliance with the provisions of 37 CFR 1.97. Accordingly, the information disclosure statement is being considered by the examiner. A signed copy of the IDS is attached hereto.

#### Response to Arguments

As Applicant's arguments appear to be solely drawn to the current Amendment file on 8/31/2007, which has not been entered, such arguments have not been considered.

As such the following rejections are maintained:

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## Claim Rejections - 35 USC § 112

The following is a quotation of the second paragraph of 35 U.S.C. 112:

The specification shall conclude with one or more claims particularly pointing out and distinctly claiming the subject matter which the applicant regards as his invention.

Claims 47-48, 54, 57 and 61-63 remain rejected under 35 U.S.C. 112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject matter which applicant regards as the invention.

Claims 47, 54 and 57 are rejected under 35 U.S.C. 112, second paragraph, as being incomplete for omitting essential steps, such omission amounting to a gap between the steps. See MPEP § 2172.01. The omitted steps are: the active of steps of determining. In the instant case, it is unclear what activity is being determined or how this activity and/or interaction is being determined.

The following is a quotation of the first paragraph of 35 U.S.C. 112:

The specification shall contain a written description of the invention, and of the manner and process of making and using it, in such full, clear, concise, and exact terms as to enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to make and use the same and shall set forth the best mode contemplated by the inventor of carrying out his invention.

Claim 61 remains rejected under 35 U.S.C. 112, first paragraph, as containing subject matter which was not described in the specification in such a way as to reasonably convey to one skilled in the relevant art that the inventor(s), at the time the application was filed, had possession of the claimed invention. Claim 61 recites a method for determining whether an agent affects one or more Mdm2-associated, HAUSP-associated, or p53 associated biological events in a cell, comprising the steps of contacting a cell with an agent that is reactive with Mdm2 or HAUSP, as determined by the method of claim 54 or 57. Thus, the claims encompass using a genus of molecules identified in a screening assay.

The specification teaches that modulators of MDM2 and HAUSP can be easily identified by simple screening assays. For example, the specification teaches that cells (e.g., MDM2-null cells or cells comprising MDM2) can be plated onto microtiter plates, then contacted with a library of drugs (paragraph 00128). In particular, the specification teaches that modulators include, but are not limited to, proteins, polypeptides, peptides, nucleic acids (including DNA or RNA), antibodies and fragments thereof, molecules, compounds, antibiotics, drugs, an agent reactive with HAUSP that induces or upregulates HAUSP expression, an agent reactive with MDM2 that induces or

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upregulates MDM2 expression, and an agent reactive with an MDM2-HAUSP complex. Thus, while the specification contemplates a number of potential modulators which can be used a screening assay, the written description does not appear to be commensurate in scope with the claimed invention because the claims encompass a genus of compounds defined only by their function wherein the relationship between the structural features of members of the genus and said function have not been defined. In the absence of such a relationship either disclosed in the as filed application or which would have been recognized based upon information readily available to one skilled in the art, the skilled artisan would not know how to make and use compounds that lack structural definition. The fact that one could have assayed a compound of interest using the claimed assays does not overcome this defect since one would have no knowledge beforehand as to whether or not any given compound (other than those that might be particularly disclosed in an application) would fall within the scope of what is claimed. It would require undue experimentation (be an undue burden) to randomly screen undefined compounds for the claimed activity.

Although the description does not provide working examples, the description teaches a method for measuring the biochemical and binding activity of the specific MDM2-HAUSP interaction, and the person skilled in the art can understand how to use the screening method considering the common general knowledge.

To comply with the written description requirement of 35 U.S.C. § 112, first paragraph, a patent specification must describe the claimed invention in sufficient detail that one skilled in the art can reasonably conclude that the inventor had possession of the claimed invention. An applicant shows possession of the claimed invention by describing the claimed invention Edith all of its limitations using such descriptive means as words, structures, figures, diagrams, and formulas that fully set forth may be shown in a variety of ways including description of an actual reduction to practice, or by showing that the invention was Already for patenting" such as by the use of drawings or structural chemical formulas that show that the invention was complete, or describing distinguishing identifying characteristics sufficient to show that the applicant was in Possession of the claimed invention.

Claimed invention is drawn to an agent identified by the method of claims 54 or 57. However, no structural or specific functional characteristics of such an agent is provided, nor is there any indication that the artisan actually implemented the method of claims 54 or 57 so as to

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identify any agent. This situation is analogous to that of Regents of the University of California v Eli Lilly, 119 F.3d 1559, 43 USPQ2d 1398 (Fed. Cir. 1997). Because one skilled in the art would conclude that the inventors were not in possession of the claimed invention. The claim fails to comply with the written description requirement.

Therefore, No claim is allowed.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Brandon J. Fetterolf, PhD whose telephone number is (571)-272-2919. The examiner can normally be reached on Monday through Friday from 7:30 to 4:30.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Shanon Foley can be reached on 571-272-0898. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

Brandon J Fetterolf, PhD Patent Examiner Art Unit 1642

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